

IN THE SPECIFICATION

Please replace the paragraph beginning at page 2, line 15 as follows:

With respect to such a urethane (meth)acrylate oligomer, molecular designing can easily be done by changing the starting material variously depending upon the ~~particularly~~ particular purpose of its use. Especially, it is easy to change the performance by the polyol to be used. When a low molecular weight polyol having a molecular weight of about a few hundreds, is used, a hard and brittle cured product will be obtained, and when a high molecular weight polyol having a molecular weight of from a thousand to a few thousands, is employed, a flexible and tough cured product will be obtained. As such a high molecular weight polyol, a polyoxyalkylene polyol, a polyester polyol, a polycaprolactone polyol or a polycarbonate polyol is, for example, employed.

Please replace the paragraph beginning at page 7, line 15 as follows:

Specifically, it may be a polyisocyanate such as tolylene diisocyanate, diphenylmethane diisocyanate, polymethylenepolyphenylene isocyanate (crude MDI), xylylene diisocyanate, isophorone diisocyanate or hexamethylene diisocyanate, or its prepolymer-modified product, ~~nulate-modified~~ isocyanurate-modified product, urea-modified product or carbodiimide modified product. Among them, a diisocyanate having two isocyanate groups in one molecule, is preferred. These polyisocyanate compounds may be used alone or in combination as a mixture of two or more of them.

Please replace the paragraph beginning at page 9, line 26 as follows:

The reaction is preferably carried out at a temperature for a common urethane-modification reaction i.e. from 30 to 90°C. At the time of the reaction, a urethane-modification catalyst such as cobalt naphthenate, zinc naphthenate, lead 2-ethylhexanoate,

dibutyltin dilaurate, tin ~~2-ethylhexanoate~~ 2-ethylhexanoate, triethylamine or 1,4-diazabicyclo[2.2.2]octane, may be used.

Please replace the paragraph beginning at page 11, line 23 as follows:

Further, for the composition of the present invention, for the purpose of lowering the viscosity, the following ethylenically unsaturated group-containing compound may be used as a diluting agent. As such an ethylenically unsaturated group-containing compound, in addition to the above-mentioned hydroxylated (meth)acrylate compound (C) such as 2-hydroxyethyl(meth)acrylate, a (meth)acrylate compound containing no hydroxyl group, such as an alkyl (meth)acrylate such as methyl (meth)acrylate or ethyl (meth)acrylate, phenoxyethyl (meth)acrylate or ~~isobornyl~~ isobornyl (meth)acrylate, or a vinyl ether monomer such as hydroxybutyl vinyl ether, lauryl vinyl ether or 2-ethylhexyl vinyl ether, may be mentioned. The urethane (meth)acrylate oligomer of the present invention has a low viscosity by itself, and in many cases, such a diluting agent may not be required. It is preferred not to use such a diluting agent.

Please replace the paragraph beginning at page 13, line 2 as follows:

Polyol A, C: ~~Prepared~~ Is prepared by reacting propylene oxide to an initiator (a polyoxypropylene diol having a molecular weight of 700) using zinc hexacyanocobaltate-glyme complex as a catalyst, and then, deactivating the catalyst, followed by purification.

Please replace the paragraph beginning at page 13, line 7 as follows:

Polyol B: One is prepared by reacting propylene oxide to an initiator (a polyoxypropylene diol having a molecular weight of 700) using zinc hexacyanocobaltate-

glyme complex as a catalyst, then reacting ethylene oxide, then deactivating the catalyst, followed by purification. The oxyethylene group-content is 20 wt%.

Please replace the paragraph beginning at page 13, line 13 as follows:

Polyol D: ~~Prepared~~ Is prepared by reacting propylene oxide to an initiator (a polyoxypropylene triol having a molecular weight of 1,000) using zinc hexacyanocobaltate-glyme complex as a catalyst, then deactivating the catalyst, followed by purification.

Please replace the paragraph beginning at page 13, line 18 as follows:

Polyol E: ~~Prepared~~ Is prepared by reacting propylene oxide to an initiator (a polyoxypropylene diol having a molecular weight of 700) zinc hexacyanocobaltate-t-butanol complex as a catalyst, then deactivating the catalyst, followed by purification.

Please replace the paragraph beginning at page 13, line 23 as follows:

Polyol F: ~~Prepared~~ Is prepared by reacting propylene oxide to an initiator (a polyoxypropylene diol having a molecular weight of 700) using potassium hydroxide as a catalyst, followed by purification. In Table 1, the number of hydroxyl groups, the hydroxyl values V_{OH} (mgKOH/g), the total degrees of unsaturation V_{US} (meq/g) and values x ($x=(0.45/V_{OH})+0.02$) are shown.

Please replace the paragraph beginning at page 14, line 7 as follows:

To 100 parts by weight (hereinafter referred to simply as parts) (0.025 mol) of polyol A, 8.7 parts (0.05 mol) of tolylene diisocyanate (T-100, tradename, manufactured by Nippon Polyurethane Industry Co., Ltd.) ~~was~~ is added, followed by a reaction at 80°C for 4 hours in the presence of 0.01 part of dibutyltin dilaurate, to obtain a urethane prepolymer. To this

urethane prepolymer, 0.05 part of dibutyltin dilaurate and 0.05 part of hydroquinone monomethyl ether ~~were~~ is added, and 6.38 parts (0.055 mol) of 2-hydroxyethyl acrylate ~~was~~ is dropwise added at 40°C. Thereafter, a reaction ~~was~~ is carried out at 60°C until absorption by an isocyanate group at a wavelength of 2,250 cm⁻¹ in the infrared absorption spectrum ~~was~~ is no longer observed, to obtain a urethane acrylate oligomer.

Please replace the paragraph beginning at page 15, line 3 as follows:

A urethane acrylate oligomer ~~was~~ is obtained in the same manner as in Example 1 except that polyol B ~~was~~ is used instead of polyol A.

Please replace the paragraph beginning at page 15, line 7 as follows:

A urethane acrylate oligomer ~~was~~ is obtained in the same manner as in Example 1 except that instead of 100 parts of polyol A, a mixture comprising 182 parts (0.0182 mol) of polyol C and 45 parts (0.0045 mol) of polyol D, ~~was~~ is used, to obtain a urethane acrylate oligomer.

Please replace the paragraph beginning at page 15, line 13 as follows:

A urethane acrylate oligomer ~~was~~ is obtained in the same manner as in Example 1 except that instead of polyol A, polyol E ~~was~~ is used.

Please replace the paragraph beginning at page 15, line 17 as follows:

A urethane acrylate oligomer ~~was~~ is obtained in the same manner as in Example 1 except that instead of polyol A, polyol F ~~was~~ is used.

Please replace the paragraph beginning at page 15, line 21 as follows:

A urethane acrylate oligomer ~~was~~ is obtained in the same manner as in Example 1 except that instead of 100 parts of polyol A, 50 parts (0.025 mol) of polyoxytetramethylene glycol having a molecular weight of 2,000, ~~was~~ is used, to obtain a urethane acrylate oligomer.

Please replace the paragraph beginning at page 15, line 27 as follows:

Examples 1 to 6 ~~were~~ may be evaluated by the following methods. The results possible are shown in Table 2.

Please replace the paragraph beginning at page 16, line 3 as follows:

The viscosity (cP) at 25°C ~~was~~ is measured by means of an E-type rotary viscometer.

Please replace the paragraph beginning at page 16, line 7 as follows:

To 100 parts of the urethane acrylate oligomer in each of Examples 1 to 6, 3 parts of benzophenone and 1 part of 2-hydroxy-2-methyl-1-phenylpropan-1-one ~~were~~ is added and thoroughly is mixed at 60°C, to obtain a photo-curable composition. The obtained photo-curable composition ~~was~~ is coated on an OPP film (Oriented polypropylene film) bonded to a glass plate by means of an applicator of 8 mil and is irradiated for 5 seconds by a high pressure mercury lamp having an output of 80 W/cm from a height of 15 cm, to obtain a cured film.

Please replace the paragraph beginning at page 16, line 17 as follows:

The surface tackiness of the cured film ~~was~~ is evaluated by touching with a finger. Evaluation of the surface tackiness ~~was~~ is made based on the following standards. ◎: No tackiness, ○: no substantial tackiness, Δ: slight tackiness, and ×: substantial tackiness.

Please replace the paragraph beginning at page 16, line 23 as follows:

Further, the mechanical properties of the film i.e. the tensile strength (unit: kg/cm²) and the break elongation (unit: %), ~~were~~ is measured in accordance with JIS K6301.